

Instructions for Use

REF Universal Prepper 800-0161

Replacement Pad
Universal Prepper Pad 508-0159

INTENDED USE

Intended use is to hold upper or lower extremities safely and effectively, with its unique design, allowing for infinite positioning of extremity during prepping procedure. The intended users of this device are medical professionals within hospitals and surgery centers.

INSTRUCTIONS

Become familiar with patient positioning device's features before use with patient. Always practice on a nurse, physician or appropriate volunteer prior to using clinically.



Attaching Universal Prepper to Surgical Table

- 1. Attach Schure Socket XL P/N 800-0134 (sold separately) to side rail
- 2. Insert mounting post into socket locking tight
- 3. Unlock horseshoe cradle and move to desired position, turn handle to lock

Detaching Universal Prepper from Surgical Table

1. Turn clamp handle counterclockwise to loosen and remove

GENERAL SPECIFICATIONS

Device Dimensions (maximum)

- Length: 36.5" +/- 0.5" (93 cm +/- 1 cm)
- Width: 11" +/- 0.5" (28 cm +/- 1 cm)
- Depth: 10" +/- 0.5" (25 cm +/- 1 cm)
- Device Weight: 5 +/- 0.5 lbs. (2 +/- .22 kg)
- 5/8"(1.6 cm) diameter mounting post insert into socket
- Single-person installation
- Twist lock/Release Handle

COMPONENT OVERVIEW

Universal Prepper is a surgical table accessory that holds upper or lower extremities safely and effectively. Unique pivoting design allows for infinite positioning of extremity.

IFU-800-0161 REV 4.00 Latest Revision: 2023-09 **Replacement Pad:** Universal Prepper Pad 508-0159

Other required product for use:

800-0134 Schure Socket XL (sold separately)

US: 0.374" x 1.122" (9.5 mm x 28.5 mm) PN# 800-0134

Denyer: 0.236" x 1.496" (6 mm x 38 mm) PN# 800-0134-DEN **Europe**: 0.394" x 0.984" (10 mm x 25 mm) PN# 800-0134-EU **Eschmann (UK)**: 0.236" x 1.260" (6 mm x 32 mm) PN# 800-0134-UK

Japan: 0.354" x 1.260" (9 mm x 32 mm) PN# 800-0134-JPN Swiss: 0.394" x 1.181" (10 mm x 30 mm) PN# 800-0134-SWISS

GENERAL INFORMATION

• Product not made with Natural Rubber Latex

- Device supports 500 lb. (227 kg) proportional patient load (6'4" (193 cm) tall patient per 99% human body model)
- Product warranty covers product from manufacturing defects for period of 2 years
- If damaged in shipping, call Customer Service at (888) 724-8763 or (781) 982-7000 to obtain Return Material Authorization number. For product warranty issues, contact Customer Service.
- CE marked medical device according to MDR (EU) 2017/745
- Product is maintenance-free, check product condition before next use
- Life of device is 5 years under normal use
- Store device between -4°F to +86°F (-20°C to 30°C)

DISPOSAL

- General Prevent infection by cleaning and disinfecting product before disposal
- Packaging Dispose packaging material via household waste according to national requirements
- SchureMed accepts back used or retired products or dispose of product in accordance with national requirements



PRODUCT USE WARNINGS

WARNING! Maximum load should not exceed appropriate proportion of a patient weighing 500 lbs. (227 kg). Use care with low-maximum load capacity surgical tables that accessory rails are not overloaded.

WARNING! Hazards result from incorrect use. Strictly follow Instructions for Use with your Operating Table system.

WARNING! Surgical table load capacities may be less. Never overload a surgical table. Device is intended for mounting on side rail of surgical table only.



WARNING! Do not reuse device if there are obvious signs of damage or functional issues. Consult manufacturer before reusing.

CLEANING RECOMMENDATION

Follow current Association of periOperative Registered Nurses Journal Guidelines for proper cleaning and disinfection procedures.



WARNING!

Adhere to standards for blood-borne pathogens from Occupational Safety and Health Administration. Use recommended protective clothing, gloves, masks and eye protection to clean accessory.

CAUTION

Strictly read/follow manufacturer's directions for cleaning fluids. DO NOT use cleaners containing phenolics.

- 1. Remove major contaminants from accessory with disposable materials. Follow appropriate bio-hazard waste disposal procedures.
- 2. Apply cleaning fluid liberally to entire accessory and wipe with clean, lint-free cloth until all moisture and cleaning fluid is removed from accessory
- 3. Let accessory dry

USER NOTICE

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

eIFU Language Versions

To download and print the Instructions for Use, please go to www.schuremed.com

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Symbol Glossary

Symbol	Title
***	Manufacturer
M	Date of manufacture
EC REP	Authorized Representative in the European Community
CH REP	Authorized Representative in the Swiss
	Importer
SN	Serial Number
À	Warning
MD	Medical Device
UDI	Unique Device Identifier
CE	CE Marking
8	Single Patient Use



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